PTC/SB/08a (08-03)
Approved for use through 07/31/2006 ONE 0651-0201
U.S. Patent and Trademark Office, U.S. DEPARTMENT OF COMMERCE
to a collection of information unless it contains a valid OMS control number. Under the Paperwork Reduction Act of 1995, no persons are req

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)	Application Number	
	Filing Date	
	First Named Inventor	Gereon VOGTMEIER
	Art Unit	1
	Examiner Name	•
	Attorney Docket Number	PHDE030413US

	Remove					
		Kind Code ¹			Pages,Columns,Lines where Relevant Passages or Relevan Figures Appear	
	1	4492869		1985-01-08	Suzuki, et al.	ail
	2	4845363		1989-07-04	Akai	all
	3	5929449		1999-07-27	Huang	all
	4	6292528	B1	2001-09-18	Wieczorek, et al.	all
	5	6324244	B1	2001-11-27	Lauter, et al.	all
	6	6495845	B1	2002-12-17	Tsunota, et al.	all
	7	6982423	B2	2006-01-03	Elgah	all
If you wis	h to ac	dd additional U.S. Pa	tent citatio	n information p	lease click the Add button.	Add

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		
iling Date		
First Named Inventor Gere		on VOGTMEIER
Art Unit		
Examiner Name		
Homey Docket Numb	or	PHDE030413US

Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publica Date	ition	Name of Pat of cited Docu	entee or Applicant ument	e or Applicant Delevent		Columns, Lines where ant Passages or Relevant s Appear	
	1	20020011572	A1	2002-0	1-31	Kajiwara, et a	4.	all			
If you wish	h to a	dd additional U.S. Publi	shed Ap	plication	citatio	n information	please click the Ad	d butto	n. Add		
				FOREIG	3N PAT	TENT DOCUM	IENTS		Remove		
Examiner Initial*	Cite No	Foreign Document Number ³	Countr Code ²		Kind Code4	Publication Date	Name of Patente Applicant of cited Document		where Re	or Relevant	TS
	1	WO 03/044563	wo		A1	2003-05-30	Elgali		all		
If you wish	h to a	dd additional Foreign P	atent Do	cument	citation	information p	lease click the Add	buttor	Add		
			NON	I-PATE	NT LITE	RATURE DO	CUMENTS		Remove		
Examiner Initials*	caminer Citie Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, dity and/or country where published.						Τs				
	1										

EXAMINER SIGNATURE

| Date Considered |
| EXAMINER: Initial if reference considered, whether on not citation is in conformance with MPEP 609. Draw line through a citation if not incomformance and not considered. Include copy of this form with next communication to applicant.

If you wish to add additional non-patent literature document citation information please click the Add button Add

See Kinz Codes of USPTO Patient Documents at year USPTO_CODE or MPEP 901.6. If Earls office that assed the document, by the holefar code (WIPO Standard ST.3). Sort Justinese plants from Coursels, the includes of the year of the Proprior many procecible the serial number of the plants of the Standard ST.16 if possible. If Applicant is to place a check mark here if English tanguage the installation is attacked.

Application Number Fing Date STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99) Application Number Fing Date First Number Invalidation Gereon VOGTMEIER Art Unit Examiner Name Attorney Docket Number | PHDE030413US

CERTIFICATION STATEMENT

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 37 CPR 1.97(e)(1).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patter office in a counterpart foreign application, nd, to the knowledge of the person signing the certification after making reasonable inquiry, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 155(c) more than three months prior to the filing of the information disclosure statement. Sea 7 CFR 137(c)(c)

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

Please see 37 CFR 1.97 and 1.98 to make the appropriate selection(s):

.7 None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

_			
Signature	/TML/	Date (YYYY-MM-DD)	2006-06-01
Name/Print	Thomas M. Lundin	Registration Number	48979

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested to the patient process and/or examine your submission, which may related that the patient process and/or examine your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pursuant to 5 U.S.C. 552a(m.).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uting an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2014 and 2016. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant for the control of t
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via flori of mapplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patient.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.